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Amendments to the Claims:

This claim listing will replace all prior versions and listings of claims in the application:

Claim Listing:

1-17. (Canceled)

- 18. (Currently Amended) A method for inducing an immune response in a mammal, the method comprising administering to the mammal a compound comprising a CpG dinucleotide and an immunomodulatory moiety wherein the immunomodulatory moiety is selected from the group consisting of: one or more abasic nucleoside, 1,3-propanediol linker, 3'-3' linkage and a modified base-containing nucleoside, wherein the modified base-containing nucleoside is selected from the group consisting of: inosine, 2-aminopurine, nebularine, 7-deaza-guanosine, nitropyrrole, nitroindole, deoxyuridine, 4-thiodeoxyuridine, d-isoguanosine, d-iso-5-methylcytosine and P-base; and wherein the compound has greater or reduced immunostimulatory effect than it would have if it lacked the immunomodulatory moiety.
- 19. (Original) The method according to claim 18, wherein the mammal is a human.
- 20. (Original) The method according to claim 18, wherein the administration of the compound is parenteral, oral, sublingual, transdermal, topical, intranasal, intratracheal, or intrarectal.
- 21. (Previously Presented) The method according to claim 18, wherein the compound is administered at a sufficient dosage to attain a blood level of oligonucleotide from about 0.01 micromolar to about 10 micromolar.
- 22. (Original) The method according to claim 18, wherein dosage of compound is from about 0.1mg per patient per day to about 200mg per kg body weight per day.
- 23. (Original) The method according to claim 18, wherein the compound is administered in combination with a vaccine.

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24. (Previously Presented) The method according to claim 23, further comprising administering an adjuvant.

25-26. Cancelled.

27. (Previously Presented) The method according to claim 18, wherein G is selected from the group consisting of guanosine, 7-deazaguanosine and inosine.